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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Appl. No.

: 10/517,518

Applicant

: Stefan SPERL

Filed

: July 1, 2005 (371 filing date)

TC/A.U.

: 1614

Examiner

: Not yet assigned.

Docket No.

: 2923-671

Customer No.

: 6449

Confirmation No.

: 1246

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents P. O. Box 1450 Alexandria, VA 22313-1450

Sir:

On July 1, 2005, Applicant filed this application with an Information Disclosure Statement submitting the International Search Report in the corresponding international application. We now enclose copies of the references that have been received from Applicant along with a copy of the previously filed PTO-1449 for the Examiner's convenience. Counsel does not have English translations of the foreign patent documents. English language abstracts are included with printed publications WO 02/074756, WO 0004954 and WO/0170204. An English language abstract of DE19940389 and DE10029014 is also enclosed.

In the event that any fees are due with this paper, please charge our Deposit Account No. 02-2135.

Respectfully submitted,

Bv

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SEP 16 JUL THE				Application Number	10/517,518	
INFORMA	TION DISC	LOSU	RE	Filing Date	December 3, 2004	
INFORMATION DISCLOSURE				First Named Inventor	Stefan SPERL	
				Group Art Unit		
				Examiner Name		
				Confirmation No.		
Sheet	1	of	2	Attorney Docket Number	2923-671	

FOREIGN PATENT DOCUMENTS								
Examiner Initials*	Cite No.1	Office ³ Code			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	T ⁶	
	1.	wo	02/074756	А	Pentapham AG	9/26/02		
	2.	wo	00/04954	А	Stuerzebecher et al	2/3/00		
	3.	DE	199 40 389	Α	Wilex Biotechnology GmbH	3/1/01		
	4.	DE	100 29 014	А	Univ Schiller Jena	12/20/01		
	5.	wo	01/070204	A3	Max Planck	9/27/01		
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Examiner Signature					Date Considered		- · · ·	

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹Unique citation designation number. ²See attached Kinds of U.S. Patent Documents. ³Enter Office that issued the document, by the two-letter code. ⁴For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶Applicant is to place a check mark here if English language translation is attached. AB indicates that only an English language abstract is attached.

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6 100. E				Application Number	10/517,518	
STATEMENT BY APPLICANT				Filing Date	December 3, 2004	
				First Named Inventor	Stefan SPERL	
				Group Art Unit		
				Examiner Name		
.=				Confirmation No.		
Sheet	2	of	2	Attorney Docket Number	2923-671	

NON PATENT LITERATURE DOCUMENTS							
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published					
	6.	Sturzebecher et al., "3-Amidinophenylalanine-based inhibitors of Urokinase", BIOORGANIC & MEDICINAL CHEMISTRY LETTERS, vol. 9, no. 21, November 1, 1999, pgs. 3147-3152.					
	7.	Magdolen et al., "Natural and synthetic inhibitors of the tumor-associated serine protease urokinase-type plasminogen activator", "ADVANCES IN EXPERIMENTAL MEDICINE AND BIOLOGY, vol. 477, 2000, pgs. 331-341.					
	8.	Heechung et al., "Selective Inhibition of Urokinase by Substituted Phenylguanidines:", JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY, vol. 33, no. 11, 1990, pgs. 2956-2961.					
	9.	Sperl et al., "(4-Aminomethyl)phenylguanidine derivatives as nonpeptidic highly selective inhibitors of human urokinase", PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES, vol. 97, no. 10, 9 May 2000, pgs. 5113-5118.					
	10.	Nienaber et al., "Re-engineering of human urokinase provides a system for structure-based drug design at high resolution and reveals a novel structural subsite", JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 275, no. 10, 10 March 2000, pgs. 7239-7248.					
Examiner Signature		Date Considered					

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